California Drug Recall Information

Recall Name
Mylan Institutional Recalls Hydrocodone Bitartrate and Acetaminophen Tablets, USP 10mg/500mg Manufactured by Qualitest Pharmaceuticals Due to the Potential for Oversized Tablets

Recall Date	Product Description	Recalling Firm	Recall Reason
12/20/12	Hydrocodone Bitartrate and Acetaminophen Tablets, USP 10mg/500mg NDC# 0603-3888-21	**Mylan Institutional, Inc.** Canonsburg, PA	*Potential for Super-Potency due to Oversized Tablets*
Recall Class	**Product Identification**	**Distribution**	**Affected Dates**
I	Hydrocodone Bitartrate and Acetaminophen Tablets, USP 10mg/500mg, 100 ct. Suspect Lots Recalled: • 3037841 • 3040859 • 3042573 Tablets are pink, capsule-shaped tablets with "***3600***" debossed on one side of the tablet and "***V***" on the other.	**CA**, nationwide **Note**: Mylan Institutional repackaged and distributed the product under the UDL Laboratories, Inc. (n/k/a Mylan Institutional Inc.) label	Distributed between February 20, 2012 and November 19, 2012

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

http://www.fda.gov/Safety/Recalls/ucm331218.htm